Arf-like Protein 3 (ARL3) Regulates Protein Trafficking and Ciliogenesis in Mouse Photoreceptors*

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Arf-like protein 3 (ARL3) is a ubiquitous small GTPase expressed in ciliated cells of plants and animals. Germline deletion of Arl3 in mice causes multigorgan ciliopathy reminiscent of Bardet-Biedl or Joubert syndromes. As photoreceptors are elegantly compartmentalized and have cilia, we probed the function of ARL3 (ADP-ribosylation factor (Arf)-like 3 protein) by generating rod photoreceptor-specific (prefix rod) and retina-specific (prefix ret) ARL3 deletions. In predegenerate retARL3−/− mice, lipidated phototransduction proteins showed trafficking deficiencies, consistent with the role of ARL3 as a cargo displacement factor for lipid-binding proteins. By contrast, rodARL3−/− rods and cones expressing Cre recombinase during embryonic development formed neither connecting cilia nor outer segments and degenerated rapidly. Absence of cilia infers participation of ARL3 in ciliogenesis and axoneme formation. Ciliogenesis was rescued, and degeneration was reversed in part by subretinal injection of adenovirus-associated virus particles expressing ARL3-EGFP. The conditional knock-out phenotypes permitted identification of two ARL3 functions, both in the expression of ARL3-EGFP. The conditional knock-out phenotypes by subretinal injection of adeno-associated virus particles Ciliogenesis was rescued, and degeneration was reversed in part outer segments and degenerated rapidly. Absence of cilia infers embryonic development formed neither connecting cilia nor outer segments and degenerated rapidly. Absence of cilia infers participation of ARL3 in ciliogenesis and axoneme formation. Ciliogenesis was rescued, and degeneration was reversed in part by subretinal injection of adenovirus-associated virus particles expressing ARL3-EGFP. The conditional knock-out phenotypes permitted identification of two ARL3 functions, both in the expression of ARL3-EGFP. The conditional knock-out phenotypes by subretinal injection of adeno-associated virus particles Ciliogenesis was rescued, and degeneration was reversed in part outer segments and degenerated rapidly. Absence of cilia infers embryonic development formed neither connecting cilia nor outer segments and degenerated rapidly. Absence of cilia infers participation of ARL3 in ciliogenesis and axoneme formation. Ciliogenesis was rescued, and degeneration was reversed in part by subretinal injection of adenovirus-associated virus particles expressing ARL3-EGFP. The conditional knock-out phenotypes permitted identification of two ARL3 functions, both in the expression of ARL3-EGFP. The conditional knock-out phenotypes by subretinal injection of adeno-associated virus particles Ciliogenesis was rescued, and degeneration was reversed in part outer segments and degenerated rapidly. Absence of cilia infers embryonic development formed neither connecting cilia nor outer segments and degenerated rapidly. Absence of cilia infers participation of ARL3 in ciliogenesis and axoneme formation. Ciliogenesis was rescued, and degeneration was reversed in part by subretinal injection of adenovirus-associated virus particles expressing ARL3-EGFP. The conditional knock-out phenotypes permitted identification of two ARL3 functions, both in the expression of ARL3-EGFP. The conditional knock-out phenotypes by subretinal injection of adeno-associated virus particles Ciliogenesis was rescued, and degeneration was reversed in part

Photoreceptor outer segments (OS)3 are modified primary cilia specializing in phototransduction (1). Polypeptides involved in phototransduction (2) are transmembrane proteins (e.g. rhodopsin) or peripheral membrane (PM) proteins (e.g. transducin and PDE6 (cGMP phosphodiesterase 6)). We are interested how PM proteins traffic from the inner segment (site of biosynthesis) through a narrow connecting cilium to the outer segment (site of phototransduction). PM proteins participating in signaling are rhodopsin kinase (GRK1, G protein receptor kinase 1) (3, 4), the heterotrimeric G protein transducin (Tαβγ) (5–8), and heterotetrameric cGMP phosphodiesterase (PDE6aβγγ) (9, 10). These proteins are soluble but lipidated (prenylated or acylated) for membrane attachment (11–16). Attachment of lipidated proteins peripherally facilitates two-dimensional diffusion for rapid interactions during phototransduction (17, 18).

Replacement of entire outer segments every 10 days in mice necessitates efficient trafficking of membrane-associated proteins (19, 20). Lipidated proteins associate transiently with the endoplasmic reticulum (ER) where post-translational processing occurs (21). Trafficking to the outer segment by diffusion requires solubilization factors that interact with lipid side chains, e.g. PDEΔ (also known as PrBP/Pr or PDE6Δ, a prenyl-binding protein originally thought to be a subunit of PDE6) (22, 23) and UNC119 paralogs (UNC119a and UNC119b, where UNC119 is uncoordinated 119, a human Caenorhabditis elegans homolog) (24). PDEΔ is a prenyl-binding protein that interacts with C-terminal farnesyl and geranylgeranyl lipids, whereas UNC119 is an acyl-binding protein specific for N-terminal C-12 and C-14 fatty acids. PDE6Δ null mutations in human are associated with Joubert syndrome (25), caused by impaired ciliary targeting of INPP5E (inositol polyphosphate-5-phosphatase E), an enzyme thought to mediate ciliary stabil-
lization (26). Deletion of Pde6d in mice produced retinal degeneration caused by trafficking defects of GRK1 and PDE6 (22). A missense mutation in UNC119 is associated with cone-rod dystrophy (27).

Arl3 (ADP-ribosylation factor (Arf)-like 3 protein) is a soluble, small GTPase that has been identified in all ciliated organisms (28). Mammalian Arl3 was identified as an expressed sequence tag (EST) and shown to be present in a number of human tissues and tumor cell lines (29). Cilia function was identified first in the protozoan Leishmania donovani (30). In experiments in ciliated hTert-RPE and IMCD3 cells (31), pull-downs (32–34), and crystalllography (35–38) identified Arf-like (ARL) proteins ARL2 and ARL3 as interactants of PDE8 and UNC119 (31, 38, 39). ARL3 localizes to the photoreceptor synaptic terminal, cell body, inner segment, and connecting cilium (40) and colocalizes with RP2 (retinitis pigmentosa protein 2) (41), UNC119 (42), and PDE8 (43). ARL3 GTase activity is regulated by a guanosine nucleotide exchange factor (GEF) and a GTPase-activating protein (GAP). RP2 functions as an ARL3 GAP, and ARL3b was identified recently as an ARL3 GEF enabling GTP/GDP exchange on ARL3-GDP (43). Mutations in ARL3b in human and mouse are associated with Joubert syndrome (44, 45). In vitro experiments identified ARL3-GTP as a guanosine nucleotide dissociation inhibitor displacement factor, and for simplicity it is referred to as cargo displacement factor (CDF). A germline Arl3 knock-out in mice revealed syndromic ciliopathy in that Arl3−/− mice did not survive beyond postnatal week 3 and sustained ciliopathy in kidney, liver, and pancreas in addition to photoreceptor degeneration (46).

We generated rod-specific and retina-specific Arl3 knock-outs to study defects in photoreceptor protein trafficking. In rodArl3 knock-outs, Cre recombinase, an enzyme carrying out site-specific recombination, is expressed post-ciliogenesis allowing formation of outer segments, whereas in retArl3 knock-outs, Cre is expressed during embryonic development. The results show that ER to OS trafficking of lipitated OS proteins in rodArl3−/− retina was impaired, leading to outer segment shortening and slow retinal degeneration, consistent with loss of CDF function. The retArl3−/− photoreceptors formed neither connecting cilia nor outer segment membrane, leading to protein accumulation in the inner segment and rapid degeneration. INPP5E, which localizes exclusively to the Golgi apparatus of WT photoreceptors, was significantly reduced in retArl3−/− inner segments suggesting that trafficking of INPP5E from ER-to-Golgi is ARL3-dependent.

**Experimental Procedures**

**Animals**—All procedures were approved by the University of Utah Institutional Animal Care and Use Committee and were conducted in compliance with the Guide for Care and Use of Laboratory Animals from National Institutes of Health. Mice were maintained in a 12:12-h dark/light cycle. Flp mice were used to invert the gene trap at FRT sites. Six3-Cre and iCre75 transgenic mice were used to generate retina- and rod-specific Arl3 knock-outs (Fig. 1) (47–49). A transgenic mouse expressing the EGFP-CETN2 fusion protein (JAX stock no. 008234) was used to identify centrioles with fluorescence microscopy (50).

**Generation of Arl3 Gene Knock-out Mouse**—A mouse embryonic stem cell line containing a gene trap cassette in intron 1 of the Arl3 gene was purchased from the European Mouse Mutant Cell Repository (EUCOMM, Helmholtz Zentrum München, Germany). The gene trap was flanked by antisense FRT and loxP sites facilitating trap inversion (51, 52) by Flp- and Cre-induced recombination (Fig. 1, B–D). The presence of the gene trap and the integrity of short and long arms were verified by PCR. Chimeric mice were generated by the transgenic mouse core facility, University of Utah. The WT allele was genotyped by PCR using primer pair Arl3-F1 (5′-AGACCAGTGCT-TTCCATC) and Arl3-R3 (5′-GTGGTTATGTTGTCTCATGAG), yielding a 367-bp amplicon. The presence of the gene trap was verified using primer pair Arl3-F1 and RP2-R5 (5′-CTAGACAAATCGGACAGAC), yielding a 1080-bp amplicon. Heterozygous Arl3 germline knock-out mice (Arl3GT/+ × GT/−) were outbred to Flp mice to invert the gene trap at FRT sites. Flp-FRT recombination was verified using the primer pair RP2-R5 and Eu-R2 (5′-CTTGAACCTCCTCGTTGAC), yielding an ~500-bp amplicon. Mice carrying the inverted gene trap (foxed mice) were mated with transgenic mice expressing Cre to generate conditional knock-outs. Cre-loxP recombination was checked with a Cre-specific primer set. The iCre75 transgene was identified using the primer pair iCre75-F (5′-GGATGCCACCTCTGATGAAG) and iCre75-R (5′-CACCAC-CTTCTTCTGACCCG). Six3-Cre mice were genotyped using the primer pair Six3Cre159 (5′-TGATGGCAACGAGTAGT-GATGAC) and Six3Cre160 (5′-TTCGGCTATACGTAACAGGG). Absence of the rd8 mutation was confirmed by PCR (54).

**Arl3 Antibody Generation**—Full-length recombinant Arl3 was prepared as described (41). Rabbit anti-Arl3 polyclonal antibody was prepared by Covance (LabCorp), Research Triangle Park, NC, using recombinant Arl3 as immunogen. Arl3 antibody was purified from bleeds using affinity chromatography on GST-Arl3. GST-Arl3 was prepared as described (41).

**Confocal Immunohistochemistry**—All animals were dark-adapted overnight. Retina cryosections were prepared as described (55). Sections were incubated with the following polyclonal primary antibodies: VPP (anti-rhodopsin, 1:500) (56); anti-rodon Tα (anti-transducin-α, 1:500, Santa Cruz Biotechnology); anti-M/L-opsin (1:500, Chemicon); anti-S-opsin (1:500, Chemicon); MOE (anti-rod PDE6, 1:500, Cytosignal); Tβγ (anti-transducin-βγ, 1:500, Cytosignal); Rab28 (goat, 1:200, Biorbyt); ARL3b (1:200, ProteinTech); and anti-Giantin (1:100, Abcam). Monoclonal antibodies included the following: G8 (anti-GRK1, 1:500, Santa Cruz Biotechnology); IS4 (anti-anguyluate cyclase 1, 1:500, from Dr. Kris Palczewski, Case Western Reserve); and CNGA1/3 (anti-CGMP-gated ion channel, 1:500, NeuroMab). Alexa488- or Alexa555-conjugated goat anti-rabbit, Cy3-conjugated goat anti-mouse, and Cy3-conjugated donkey anti-goat secondary antibodies were diluted 1:1000. Images were acquired using an Olympus Fluoview 1000 confocal microscope.

**Immunoblotting**—Proteins of retina lysates were separated by SDS–12.5% PAGE and transferred to a nitrocellulose membrane processed as described (55). Primary antibodies were diluted 1:500 for anti-Arl3 and 1:1000 for anti-β-actin (Sigma).
in 1× TTBS containing 5% milk. After incubation overnight at 4 °C, membranes were incubated for 1 h at room temperature with the following secondary antibodies (Odyssey): iR680 goat anti-mouse (1:5000) and iR800 goat anti-rabbit (1:3000). Membranes were incubated for 1 h at room temperature (3.4 log cd/s/m−2) before being washed with Odyssey stripping buffer at 4 °C, membranes were incubated for 1 h at room temperature.

**Statistics**—SigmaPlot12 was used for statistical analysis using Student’s t test, and the level of statistical significance was set to \( p = 0.05 \).
Results

Generation of Conditional Arl3 Knock-out Mice—Embryonic stem cells containing a β-GEO gene trap (52) in intron 1 of the mouse Arl3 gene were used to generate chimeric mice (Fig. 1A). Before injection into blastocysts, position of the gene trap in intron 1 and presence of inverse FRT and loxP sites flanking the trap were confirmed by PCR and sequencing (Fig. 1B). The gene trap contains a splice-acceptor sequence upstream of the promoterless β-Geo reporter gene that essentially converts the gene trap into an exon. The trap is predicted to truncate ARL3 after exon 1, which ends after the start codon ATG, thereby preventing expression of ARL3.

A. Scotopic ERG traces (n = 7) of P15 and 1-month (M)-old WT and homozygous knock-out (KO) mice at 1.4 log cd s m⁻². Changes in a-wave amplitudes as a function of flash intensity are shown (right panel). B. Photopic ERG traces (n = 7) of P15 and 1-month-old WT and homozygous knock-out mice at 1.4 log cd s m⁻². Changes in b-wave amplitudes as a function of flash intensity are shown (right panel). C. OptoMotry (n = 7) of rod-specific knock-outs at 1 month (top panel) and 2 months (bottom panel) of age. Mutant OKT responses are plotted relative to heterozygous and WT controls. OKTs were comparable with controls in 1-month-old homozygous knock-out mice but were extinguished at 2 months of age with statistical significance (p < 0.0001). D. Remnant homozygous knock-out cone outer segments, probed with anti-ML-opsin antibody (red), are present at 1 month (top panel) but not at 2 months (bottom panel), relative to WT littermates. Scale bar, 10 μm. E. WT and homozygous knock-out retina thickness measured by OCT at P15 (top panel) and 2 months (bottom panel). Scale bar, 200 μm. F. WT, homozygous knock-out and homozygous knock-out retina thickness as a function of age (n = 3), from P15 to P60. Retina degeneration starts at P15 in homozygous knock-out mice.
Photoreceptor ARL3

Germline transmission did not produce live Arl3<sup>GT/GT</sup> mice confirming embryonic or early postnatal embryonic lethality of Arl3<sup>−/−</sup> mice (46). In several Arl3<sup>GT/GT+</sup> matings, only 2 Arl3<sup>GT/GT</sup> mice (of 150) were born but did not survive (results not shown).

Arl3<sup>GT/GT+</sup> mice were mated with transgenic mice expressing flippase recombinase (flp), causing inversion of the gene trap (Fig. 1C). Inversion of the trap renders the gene trap inactive (no effect on Arl3 expression). Cre-loxP recombination was initiated with two transgenic lines, iCre75 mice expressing Cre recombinase (Cre) in rods (49, 56) or Six3-Cre mice expressing Cre in retinal progenitors (47). Cre-loxP recombination with iCre75 at postnatal day 7 (P7) and later inverted the trap only in rods to generate conditional rod knock-outs termed rod<sup>Arl3<sup>−/−</sup></sup>, whereas Cre-loxP recombination with Six3-Cre inverted the trap in all retinal progenitors generating conditional retina knock-outs termed ret<sup>Arl3<sup>−/−</sup></sup> (Fig. 1D). Genotyping of offspring (Fig. 1E) and Western blot results (Fig. 1F) confirmed the generation of each conditional knock-out line. The rd8 mutation of the Crb1 gene (54) was detected in the F1 mutant mouse line but was subsequently eliminated by successive matings with rd8-free C57BL6/J mice.

Immunohistochemistry with available Arl3 antibodies was unsuccessful. To show the distribution of Arl3 unambiguously, we generated a self-complementary adeno-associated virus (scAAV2/8) expressing the fusion protein, Arl3-EGFP. Subretinal injection of scAAV-ARL3-EGFP revealed localization of Arl3, presumably in the GDP-bound form, throughout the outer nuclear layer (ONL), inner segment (IS), and connecting cilium (CC) but not the outer segment (Fig. 1G). Arl3 is likely in the GDP (inactive) form in the inner segments because its activating protein (GAP), RP2, is present in the inner segment cell membrane (41) and its GEF (which activates GDP/GTP exchange) is located at the connecting cilium and outer segment. Arl3-GTP is needed only at the ciliary ridge to unload cargo (see below). Enlargement of the IS/OS junction (Fig. 1G, right panel) reveals that Arl3 is enriched in the CC/BB area.

rod<sup>Arl3<sup>−/−</sup></sup> Photoreceptors Degenerate with RP-like Phenotype—During retina development, the mother cilium docks to the cortex of the rod IS around P5–6 to engage in ciliogenesis and axoneme formation (59). We initiated rod deletion of Arl3 by crossing Arl3<sup>flox/flox</sup> mice with iCre75 transgenic mice expressing Cre at P7 and later (49). Scotopic ERG recordings of P15 rod<sup>Arl3<sup>−/−</sup></sup> mice at various intensities (Fig. 2A) revealed nearly normal responses at P15 but reduced rod a- and b-wave amplitudes at 1 month of age. The presence of rod responses demonstrates that postnatal development, including ciliogenesis, occurred normally. The rod<sup>Arl3<sup>−/−</sup></sup> photopic responses (Fig. 2B) were robust and comparable with WT at P15 but started to decline (15–20%) at 1 month of age, a phenotype resembling RP. Visual function was further tested by Optomotry, a technique that enables the rapid screening of functional vision using the OKT response. At 1 month of age, rod<sup>Arl3<sup>−/−</sup></sup> OKT was normal but extinct at 2 months (Fig. 2C).

The normal response at 1 month is most likely due to survival of cones (Fig. 2D, right panel). At 2 months of age, rods and cones were degenerated (Fig. 2, D and F), and the OKT response was extinguished. By optical coherence tomography (OCT), rod<sup>Arl3<sup>−/−</sup></sup> retina thickness declined at P15 (Fig. 2, E and F) and approached 160 μm at P50 (Fig. 2F). Heterozygous rod<sup>Arl3<sup>−/+</sup></sup> retina thickness (~260 μm) was identical to WT suggesting haplosufficiency (Fig. 2F). In mice expressing iCre75 on the WT background, retina thickness was unaffected over the first 60 postnatal days (56).

Arl3 Regulates Trafficking of Lipidated Peripheral Proteins—Based on ERG (Fig. 2, A and B), OKT (Fig. 2C), and retinal thickness (Fig. 2, E and F), rod<sup>Arl3<sup>−/−</sup></sup> retina presents a very early stage of degeneration at P15. ONL thickness (Fig. 3A) and rod outer segment length (Fig. 3B) are only slightly reduced, and connecting cilium length was normal (Fig. 3, C and D).
Connecting cilium, mother centriole (basal body), and daughter centriole were identified by expression of transgenic EGFP-CETN2 (Fig. 3D), a well characterized centriole (50) and connecting cilium marker (60). Transmembrane proteins, e.g. rhodopsin (Fig. 3E), rod/cone cGMP-gated channel α-subunits (CNGA1/3) (Fig. 3F), guanylate cyclase 1 (Fig. 3G), and cone...
Six3-Cre-mediated recombination is known to be delayed in the retina periphery (60, 62). We therefore tested CC/OS formation by rhodopsin immunohistochemistry in the presence of EGFP-CETN2 at P10 and P15, the times when ERG and OKT showed significant visual responses. The results show normal OS in the peripheral retArl3−/− retina, comparable with heterozygous controls, although degeneration in the central retina was far advanced (Fig. 5).

**retArl3−/− Rod and Cone Photoreceptors Do Not Form Outer Segments**—OCT of retArl3−/− animals showed a decrease in retina thickness to 200 µm as early as P15, further decreasing to ~160 µm at P30 (Fig. 6, A and B). ONL thickness is slightly reduced at P10, and active degeneration ensues at P15 (Fig. 6C). In control P10 and P15 retArl3−/− retinas, connecting cilia and outer segments are present, and functional basal bodies, connecting cilia and axonemes, were not formed (Fig. 6, D and E, right panels). Rhodopsin mislocalized throughout the ONL and IS (Fig. 6, F and G) similarly as observed in P10 retKif3a−/− retina lacking connecting cilia and outer segments (60). Trafficking of lipidated proteins (PDE6) (Fig. 6, H and I), rod Tα (Fig. 6, J and K), and GRK1 (data not shown) was also impaired as outer segment disks are absent. Pigments M/L- and S-opsin distributed throughout the retArl3−/− cones (Fig. 6, L–O). At P15, rod and cone photoreceptors degenerate massively (Fig. 6, G, I, K, M, and O, right panels). In contrast to phototransduction proteins, the ARL3 GEF ARL13b, which has an N-terminal palmitoylation motif and in C. elegans is palmitoylated (63), did not mislocalize in the retArl3−/− ONL (Fig. 6, P and Q).

**ARL13b Localization in the Mutant Retina Is Restricted to the Apical IS and Is Presumably Membrane-Associated.** The absence of outer segments as early as P10 in central retArl3−/− photoreceptors suggests that ARL3, a known microtubule-interacting protein, may regulate a late stage of ciliogenesis or intraflagellar transport.

**retArl3−/− Versus retArl3+/− Photoreceptor Degeneration**—At P15, photoreceptor degeneration in retArl3−/− mice proceeds faster than in retArl3+/− mutants due to embryonic expression of Cre and lack of outer segments in the central retina (compare Figs. 3, I–L, and 6, E–Q). At 1 month, retArl3−/− and retArl3+/− photoreceptors degenerate rapidly with profound accumulation of PDE6, GRK1, and rod Tα in the remaining inner segments and ONL (Fig. 7, A–C). One-month-old retArl3−/− and retArl3+/− retinas reveal only 3–4 or 5–6 rows of ONL nuclei (Fig. 7, A–C, and E), respectively, and at 2 months the mutant retinas are both nearly completely degenerated (Fig. 7F).

**Rescue of Ciliogenesis and Protein Trafficking in retArl3−/− Photoreceptors**—AAV particles expressing ARL3-EGFP were injected into the subretinal space of P15 WT and retArl3−/− mice. Retinas harvested at 2 months post-injection (Fig. 8, A–D) show significant rescue of the ONL layer (4–5 rows of nuclei in treated versus 1 row in untreated retinas) (Fig. 8, A–D, middle and right panels, E). Moreover, treated central retArl3−/− photoreceptors developed inner and outer segments, demonstrating rescue of ciliogenesis. Although outer

pigments (M/L- and S-opsin, results not shown), trafficked normally and were unaffected by absence of ARL3. Farnesylated rhodopsin kinase (GRK1) (Fig. 3H), prenylated rod PDE6 (Fig. 3I), acylated rod Tα (Fig. 3J), farnesylated Tβγ (Fig. 3K), and Rab28 (Fig. 3L) mistrafficked to the inner segments, the ONL, and synaptic terminals. As a consequence, lipidated proteins accumulated in the ONL and inner segments of rodArl3−/− photoreceptors as inactivation of the Arl3 gene by Cre-mediated recombination increased.

**Retina-specific Deletion of ARL3**—Six3-Cre-mediated recombination is expected to disable ARL3 expression as early as embryonic day 10 (47), before photoreceptor differentiation and ciliogenesis. Unexpectedly, retArl3−/− scotopic and photopic ERG traces were recordable at P15 and 1 month (Fig. 4, A and B). The scotopic a-waves were suppressed at P15 and highly reduced at 1 month using flash intensities of −1.63 log cd m−2 and higher (Fig. 4A, right panel). Photopic b-waves were significantly reduced in P15 and 1-month-old animals to about 35% of normal ERG traces (Fig. 4B, right panel). The retArl3−/− OKT response was 25% reduced at 1 month and nearly extinguished at 2 months (Fig. 4C). Significant visual activity in retArl3−/− animals at 1 month may be due to survival mutant cones in the peripheral retina (Fig. 4D, right panel). ML-opsin-labeled cone outer segments were clearly present in the retArl3−/− retina periphery, whereas OS were absent in the central retArl3−/− retina. At 2 months, the retArl3−/− retina was degenerated and unable to respond to light (Fig. 4D), a phenotype resembling Leber congenital amaurosis (61).

**FIGURE 5.** Fluorescent immunohistochemistry of whole retina sections of WT and retina-specific Arl3 knock-out retinas. **A–D,** sections were labeled with anti-rhodopsin (red) and contrasted with DAPI (blue) to reveal nuclei, with enlargements of the central and peripheral retina. **Scale bar,** 100 µm. **E,** ONL thickness (n = 3) as the function of the distance from the optic nerve head (ONH).
segments were shorter in treated retinas, outer segment proteins trafficked normally and did not mislocalize (Fig. 8, A–D, middle panels).

**FIGURE 6. Mistaflcicking and degeneration in** \( ^{−/−} \text{Arfl3} \) **central photoreceptors.** A, OCT of WT (left panel) and \( ^{−/−} \text{Arfl3} \) (right panel) retinas at P15 (top panel) and 2 months (bottom panel). B, retina thickness measured by OCT as a function of age (n = 3). C, ONL thickness of WT (black bars) and mutant (white bars) mice at P10 and P15 (p < 0.001; n = 3). **, p < 0.001. D–Q, retina immunohistochemistry at P10 (left panel) and P15 (right panel). Within each time point, heterozygote controls (left panel) are compared with knock-outs (right panel). Panels show EGFP-CETN2 fluorescence (D and E) or labeling with antibodies (red) directed against rhodopsin (F and G), PDE6 (H and I), rod T α (J and K), ML-opsin (L and M), S-opsin (N and O), and ARL13b (P and Q). Left quarter of each panel show DAPI (blue) to identify extent of nuclear layer. Scale bar, 10 μm.

**Trafillc of INPP5E to the Golgi—** INPP5E is present in the cilia of hTert-RPE cells, cilia of mouse kidneys and cerebellum (64), and axonemes of mouse embryonic fibroblasts (26).
INPP5E is farnesylated and was suggested to travel to the OS in a PDEδ-dependent manner (25). In WT (data not shown) and retδ+/− mouse rods, however, INPP5E localizes to the Golgi of inner segments and is absent in the outer segment (Fig. 9, A and E). At P10 when OS are forming, INPP5E is present in the inner segment proximal to centrioles, and rhodopsin localizes exclusively to the OS (Fig. 9, A and B, and enlargements). At P15 when OS are more developed, separation of rhodopsin and INPP5E fluorescence is most evident (Fig. 9, E and F). The enlargements reveal that centrioles and connecting cilia, identified by EGFP-CETN2, do not colocalize with INPP5E (Fig. 9E, bottom panel). In retδ−/− animals at P10 (Fig. 9, C and D) and P15 (Fig. 9, G and H), rhodopsin mislocalizes to the inner segments as outer segments are absent. INPP5E fluorescence of the retδ−/− IS appears significantly reduced (Fig. 9, C and G) compared with retδ+/− IS (Fig. 9, A and E), suggesting that INPP5E trafficking from ER-to-Golgi may in part be ARL3/PDEδ-dependent.

We generated an EGFP-INPP5E expression construct to avoid misinterpretation due to antibody artifacts. Neonatal electroporation of the construct and colabeling with a Golgi marker, anti-giantin, demonstrates that INPP5E distributes to the ER surrounding the nucleus and Golgi apparatus (Fig. 9, I and J), and it is excluded from WT and mutant outer segments (Fig. 9, M and N). In retδ−/− retina (Fig. 9, K and L), anti-giantin-labeled Golgi is retained in the degenerating ONL, and rhodopsin, unable to traffic, colocalizes with INPP5E (Fig. 9, O and P). The results show that farnesylated INPP5E traffics to the Golgi and not the outer segment of both WT and mutant photoreceptors.

Discussion

ARL3 is a small GTPase colocalizing with microtubules in HeLa cells and brain (40). The best-known function of ARL3, in the GTP-bound form, is that of a cargo displacement factor that evicts lipidated cargo from its lipid-binding protein (37, 38, 65). ARL3-dependent lipid-binding proteins are PDEδ and UNC119 featuring immunoglobulin-like β-sandwich structures that can accommodate lipid side chains with high affinity. However, germline Arl3−/− mice exhibited abnormal development of renal, hepatic, and pancreatic epithelial tubule structures, implicating ARL3 in ciliogenesis (46) and syndromic ciliopathy. To explore the role of ARL3 in photoreceptors, we generated rod- and retina-specific Arl3 knockouts. An advantage of the rod-specific knock-out is that Cre recombinase is expressed relatively late (>7), after ciliogenesis has occurred, allowing rodδ−/− rod outer segments to form normally. We observed significant mis trafficking of lipidated proteins as early as P15 (Fig. 3, H–L, right panels), signaling the onset of an RP-like photoreceptor degeneration. Rhodopsin and other transmembrane proteins are unaffected by loss of ARL3. In the WT scenario, lipidated cargo, docked to the ER membrane post-biosynthesis, is extracted by its lipid-binding protein (PDEδ or UNC119) and forms a soluble complex (Fig. 10A). The complex interacts with its CDF (ARL3-GTP), expelling cargo from its binding site and delivering it to the destination membrane. The phenotype of the rod-specific Arl3 knock-out is consistent with loss of CDF function (Fig. 10B). In the absence of ARL3, lipidated cargo is retained and not delivered to the destination membrane, leading to accumulation of lipidated proteins in the IS and ONL (Figs. 3, H–L, and 10B). Non-delivery of PDE6 to the outer segment poses significant problems by upsetting the PDE6/guanylate cyclase cGMP regulatory system and shifting free cGMP to higher levels. As a result, rods degenerate with an RP-like phenotype that eventually affects cones as well (66).

Homozygous Rp2 null alleles in humans cause severe X-linked RP (XLGRP), whereas Rp2 deletion in mice promotes a slowly progressing retinal degeneration (41, 67). We proposed that an abundance of dominantly active ARL3-GTP, stabilized by negligible intrinsic GTPase activity, attenuates the ability of PDEδ to interact with prenylated cargo (41). Interaction of ARL3-GTP with its PDEδ complex constricts the lipid-binding site and impairs the binding of cargo (alternatively, for UNC119, the binding pocket widens). Failure to extract cargo from the ER steadily lowers the concentration of lipidated proteins in the OS, initiating a slowly progressing photoreceptor degeneration. The rodδ−/− trafficking defects are distinct from the Rp2−/− mouse defects where ARL3 is locked in the GTP-bound form thereby impeding extraction of PDE6 and GRK1 from the ER and leading to the degradation of the lipidated proteins (41). Over-
expression of dominantly active ARL3 in *C. elegans* and *Leishmania* also causes ciliogenesis defects (30, 68), but mistrafficking of lipidated proteins has not been invoked as causal.

The severe ciliopathy phenotype of *Arl3* germline knockouts prompted us to investigate the consequence of ARL3 deletion during embryonic and early postnatal development, before the onset of ciliogenesis. Using transgenic EGFP-CETN2 as centriole and connecting cilium marker, we observed that connecting cilia (transition zones) and axonemes were undetectable in the central P10 *ret*<sup>Arl3<sup>−/−</sup> retinas, and the outer segments never formed (Figs. 5 and 6, *D* and *E*). As an immediate consequence, transmembrane and peripheral membrane outer segment polypeptides accumulated in the inner segment (Fig. 6, *D*–*Q*). Although severe retinal degeneration was obvious at

![Graphical representation](image-url)
P10, ciliogenesis in central retinal photoreceptors could be ameliorated by viral expression of ARL3 tagged with EGFP (Fig. 8). Although rescue of degeneration was only partial, inner segment and shortened outer segments were present in treated retinas demonstrating rescue of ciliogenesis and protein trafficking. Compartmentalization of ARL3 in the IS/basal body area and of its GEF, ARL13b, in the OS suggests that ARL3 must be switched into its GTP-bound active conformation to enable ciliogenesis. Accordingly, mutations in the human ARL13b gene disabling GEF activity on ARL3 and germline knock-out of the mouse Arl13b gene cause Joubert syndrome (44, 45).

The \(^{\text{ret} \text{Arl3}^{-/-}}\) phenotype is strikingly similar to a retina-specific knock-out of KIF3A (obligatory subunit of the heterotrimeric kinesin-2, KIF3) and IFT88 (particle of the IFT-B complex) (60). KIF3 and IFT88 are responsible for IFT of tubulin and axoneme building blocks. In \(^{\text{ret} \text{Kif3a}^{-/-}}\) rod and cone photoreceptors, ciliogenesis was disabled as early as P6 in the central retina, with a delay in the peripheral retina. A link between
ARL3 and intraflagellar transport was observed in *Leishmania* and *C. elegans*. In *Leishmania*, ARL3 is required for flagella formation (69), and in *C. elegans* ARL13b and ARL3 regulate IFT through a tubulin deacetylase (HDAC6) pathway (28, 68). In *Arl3* mutant worms, which have no GEF activity and presumably lack ARL3-GTP, IFTA, and IFTB, particles dissociate and disrupt IFT (28) suggesting that ARL3-GTP may be required for assembly of IFT complexes.

Finally, we investigated the trafficking of INPP5E in photoreceptors, dependence on ARL3/PDE6, and as a phosphatidylinositol-5'-phosphatase. Mutations in the human *INPP5E* gene are linked to Joubert and MORM syndromes (26, 64), and depletion of INPP5E in the adult mouse by tamoxifen induction causes rapid photoreceptor degeneration (26), suggesting a key role for phosphoinositide metabolism in ciliary maintenance. INPP5E, originally cloned from Golgi membranes, was shown to be predominantly located to the COS-7 (70). Among ciliated cells, INPP5E colocalized with acetylated α-tubulin in hTert-RPE cells, localized to cilia of P10 mouse kidneys and P5 cerebellum (64), and is concentrated in the axoneme of mouse embryonic fibroblasts (26). In IMCD3 cells, targeting to Golgi, to the OS was proposed to be PDE6/ARL3-dependent, whereas targeting to cilia was ARL13b-dependent (71). Absence of PDE6 prevented delivery of INPP5E to mouse embryonic fibroblast cilia, and a *PDE6D* null allele was linked to Joubert syndrome (25).

INPP5E was absent in mouse photoreceptor outer segments and apical inner segments and instead localized to the proximal inner segment and Golgi (Fig. 9, A–H). This is surprising, as INPP5E localizes along the axoneme of ciliated cells (26). Furthermore, PDE6-mediated trafficking of INPP5E to the OS was suggested to be impaired in patients carrying a PDE6D null allele (25). Absence of INPP5E in the OS and its localization to the Golgi implies a critical role for INPP5E in the latter organelle. INPP5E levels in the *rod*/*Arl3* inner segment/Golgi were reduced, suggesting that ER-to-Golgi trafficking of INPP5E may be ARL3/PDE6-dependent. ARL13b, localizing exclusively to WT and *rodArl3* OS (Fig. 6, P and Q), was suggested to target INPP5E to the OS (71). Distinct localizations of ARL13b in the OS and INPP5E in the Golgi discount this putative pathway.

The physiological role of INPP5E in the photoreceptor Golgi is intriguing as the Golgi is an organelle in which phosphatidylinositol 4-phosphate (PI4P) is enriched (72–74). PI4P could be generated by INPP5E-mediated hydrolysis of 5-phosphate bound to the inositol ring in phosphatidylinositol 4,5-bisphosphate or by 4'-phosphorylation of phosphatidylinositol by phosphatidylinositol 4-kinase. Interestingly, phosphatidylinositol 4-kinase knockdown abrogated Golgi to plasma membrane trafficking in mammalian cells (75) suggesting a key role for PI4P in vesicular trafficking. Reduction of INPP5E in the Golgi could deplete PI4P and impair Golgi-to-OS trafficking of membrane proteins, including rhodopsin, a protein that is essential for OS elaboration and vision.

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References


